Selektywny Niedobór IgM
Kod Orpha: 331235 Kod OMIM:

Opis choroby *

Definicja
A rare primary immunodeficiency characterized by recurrent and/or invasive bacterial, viral, and fungal infections, associated with low to absent blood IgM levels, while IgG, IgG subclasses, and IgA levels, as well as IgG antibody response to vaccinations, are normal. Patients may also present allergic diatheses, and the prevalence of autoimmune diseases is increased.
Dane

Klasyfikacja
Choroba

Synonimy
Selective immunoglobulin M deficiency
Selektywny Niedobór immunoglobuliny M

Kod ORPHA
331235

Kod OMIM
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Kod ICD10
D80.4

Kod ICD11
4A01.04

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.